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NEWS 3 SEP 09 ACD predicted properties enhanced in REGISTRY/ZREGISTRY  
NEWS 4 OCT 03 MATHDI removed from STN  
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NEWS 6 OCT 13 New CAS Information Use Policies Effective October 17, 2005  
NEWS 7 OCT 17 STN(R) AnaVist(TM), Version 1.01, allows the export/download  
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NEWS 11 NOV 14 CA/CAPLUS - Expanded coverage of German academic research  
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=> index bioscience

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INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, ANTE, AQUALINE,  
AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS,  
BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB,  
CROPU, DDFB, DDFU, DGENE, DISSABS, ...' ENTERED AT 11:27:52 ON 07 DEC 2005

74 FILES IN THE FILE LIST IN STNINDEX

Enter SET DETAIL ON to see search term postings or to view  
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=> (fluorescent or chromoprotein) and emission and maximum

23 FILE AGRICOLA

96 FILE ANABSTR  
 1 FILE ANTE  
 9 FILE AQUALINE  
 53 FILE AQUASCI  
 18 FILE BIOBUSINESS  
 46 FILE BIOENG  
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 70 FILE BIOTECHABS  
 70 FILE BIOTECHDS  
 207 FILE BIOTECHNO  
 64 FILE CABA  
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 7 FILE CEABA-VTB  
 7 FILE CEN  
 1 FILE CIN  
 1 FILE CONFSCI  
 8 FILE CROPU  
 9 FILE DDFU  
 594 FILE DGENE  
 84 FILE DISSABS

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 5 FILE EMBAL  
 589 FILE EMBASE  
 254 FILE ESBIODASE  
 9 FILE FEDRIP  
 7 FILE FROSTI  
 16 FILE FSTA  
 2 FILE HEALSAFE  
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 1 FILE KOSMET  
 168 FILE LIFESCI  
 774 FILE MEDLINE  
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 19 FILE OCEAN  
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 22952 FILE USPATFULL  
 2659 FILE USPAT2  
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 21 FILE WATER  
 452 FILE WPIDS  
 1 FILE WPIFV  
 452 FILE WPINDEX

51 FILES HAVE ONE OR MORE ANSWERS, 74 FILES SEARCHED IN STNINDEX

L1 QUE (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM

=> d rank

F1	22952	USPATFULL
F2	2659	USPAT2
F3	2381	CAPLUS
F4	878	SCISEARCH
F5	824	BIOSIS
F6	774	MEDLINE

F7	618	IFIPAT
F8	594	DGENE
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F10	452	WPIDS
F11	452	WPINDEX
F12	361	PASCAL
F13	254	ESBIOBASE
F14	228	TOXCENTER
F15	207	BIOTECHNO
F16	197	PROMT
F17	168	LIFESCI
F18	135	JICST-EPLUS
F19	96	ANABSTR
F20	84	DISSABS
F21	72	DRUGU
F22	70	BIOTECHABS
F23	70	BIOTECHDS
F24	64	CABA
F25	53	AQUASCI
F26	46	BIOENG
F27	43	CANCERLIT
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F41	7	CEN
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F43	5	EMBAL
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F45	2	HEALSAFE
F46	2	VETU
F47	1	ANTE
F48	1	CIN
F49	1	CONFSCI
F50	1	KOSMET
F51	1	WPIFV

=> file biosis medline caplus  
COST IN U.S. DOLLARS

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SINCE FILE ENTRY	TOTAL SESSION
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=> (fluorescent or chromoprotein) and emission and maximum and nm  
L2 2641 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND NM

=> (fluorescent or chromoprotein) and emission and maximum and "far red shifted"  
L3 1 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND  
"FAR RED SHIFTED"

=> d ab bib

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN  
AB Nucleic acid compns. encoding Stichodactylidaen **chromoproteins**  
and **fluorescent** mutants thereof, as well as the polypeptide  
compns. encoded by the same, are provided. The proteins of interest are  
proteins that are colored and/or **fluorescent**, where this feature  
arises from the interaction of two or more residues of the protein. Also  
of interest are proteins that are substantially similar to, or mutants of,  
the above specific proteins, including non-aggregating mutants and mutants  
with modulated oligomerization characteristics as compared to wild type.  
Thus, the two wild-type **chromoprotein** isoforms from Heteractis  
crispa exhibit a strong **emission max.** at  
.apprx.580-640 nm. Site-specific mutagenesis of the Cys-148 residue to  
serine dramatically increases the quantum yield of red fluorescence as  
compared to the wild-type protein, and further random mutagenesis (A2S,  
T36A, C143S, L173H, P201L, K204E) yielded an even brighter mutant. A  
single mutation, L126H, may be responsible for modifying the oligomeric  
state of the protein from tetrameric to dimeric. Also provided are  
fragments of the nucleic acids and the peptides encoded thereby, as well  
as antibodies to the subject proteins and transgenic cells and organisms.  
The subject protein and nucleic acid compns. find use in a variety of  
different applications. Finally, kits for use in such applications, e.g.,  
that include the subject nucleic acid compns., are provided.

AN 2002:293685 CAPLUS

DN 136:321045

TI **Far red-shifted fluorescent**  
proteins and their variants from Heteractis crispis

IN Lukyanov, Sergey Anatolievich; Fradkov, Arcady Fedorovich; Lukyanov,  
Konstantin Anatolievich; Gurskaya, Nadezda Georgievna

PA Clontech Laboratories, Inc., USA

SO PCT Int. Appl., 81 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 17

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	----	-----	-----	-----
PI	WO 2002030965	A2	20020418	WO 2001-US32080	20011012
	WO 2002030965	A3	20030605		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,				
	CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,				
	GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,				
	LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL,				
	PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,				
	US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG,				
	KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR,				
	IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN,				
	GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2425318	AA	20020418	CA 2001-2425318	20011012
	AU 2002011722	A5	20020422	AU 2002-11722	20011012
	EP 1334122	A2	20030813	EP 2001-979797	20011012
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,				
	IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2004532606	T2	20041028	JP 2002-534350	20011012
PRAI	US 2000-240018P	P	20001012		
	US 2001-306131P	P	20010716		
	WO 2001-US32080	W	20011012		

=> (fluorescent or chromoprotein) and emission and maximum and " red shifted"  
L4 124 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND "  
RED SHIFTED"

=> dup remove  
ENTER L# LIST OR (END):14  
PROCESSING COMPLETED FOR L4  
L5 82 DUP REMOVE L4 (42 DUPLICATES REMOVED)

=> 15 and nm  
L6 56 L5 AND NM

=> dup remove  
ENTER L# LIST OR (END):16  
PROCESSING COMPLETED FOR L6  
L7 56 DUP REMOVE L6 (0 DUPLICATES REMOVED)

=> 16 and nucleic with acid  
L8 1 L6 AND NUCLEIC WITH ACID

=> d ab bib

L8 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN  
AB **Nucleic acid** compns. encoding Stichodactylidaen  
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DN 136:321045

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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

CA 2425318 AA 20020418 CA 2001-2425318 20011012

AU 2002011722 A5 20020422 AU 2002-11722 20011012

EP 1334122 A2 20030813 EP 2001-979797 20011012

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

JP 2004532606 T2 20041028 JP 2002-534350 20011012

PRAI US 2000-240018P P 20001012

US 2001-306131P P 20010716

WO 2001-US32080 W 20011012

=> 16 and" nucleic acid"

L9 1 L6 AND" NUCLEIC ACID"

=> d ab bib

L9 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN

AB **Nucleic acid** compns. encoding Stichodactylidaen **chromoproteins** and **fluorescent** mutants thereof, as well as the polypeptide compns. encoded by the same, are provided. The proteins of interest are proteins that are colored and/or **fluorescent**, where this feature arises from the interaction of two or more residues of the protein. Also of interest are proteins that are substantially similar to, or mutants of, the above specific proteins, including non-aggregating mutants and mutants with modulated oligomerization characteristics as compared to wild type. Thus, the two wild-type **chromoprotein** isoforms from *Heteractis crispa* exhibit a strong **emission max.** at .apprx.580-640 nm. Site-specific mutagenesis of the Cys-148 residue to serine dramatically increases the quantum yield of red fluorescence as compared to the wild-type protein, and further random mutagenesis (A2S, T36A, C143S, L173H, P201L, K204E) yielded an even brighter mutant. A single mutation, L126H, may be responsible for modifying the oligomeric state of the protein from tetrameric to dimeric. Also provided are fragments of the **nucleic acids** and the peptides encoded thereby, as well as antibodies to the subject proteins and transgenic cells and organisms. The subject protein and **nucleic acid** compns. find use in a variety of different applications. Finally, kits for use in such applications, e.g., that include the subject **nucleic acid** compns., are provided.

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TI Far **red-shifted fluorescent** proteins and their variants from *Heteractis crispis*

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PRAI	US 2000-240018P	P	20001012		
	US 2001-306131P	P	20010716		
	WO 2001-US32080	W	20011012		

=> 16 and anthozoan

L10 0 L6 AND ANTHOZOAN

=> 16 and polypeptide

L11 1 L6 AND POLYPEPTIDE

=> 15 and anthozoan

L12 0 L5 AND ANTHOZOAN

=> (fluorescent or chromoprotein) and emission and maximum and " red shifted" and anthozoan

L13 0 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND " RED SHIFTED" AND ANTHOZOAN

=> d 16 ti 1-20

L6 ANSWER 1 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
TI Two-photon excitation and **emission** spectra of the green **fluorescent** protein variants ECFP, EGFP and EYFP.

L6 ANSWER 2 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
TI The two photon cross section and spectrum of 6MAP, a **fluorescent** adenosine analog.

L6 ANSWER 3 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
TI Crystallization and preliminary X-ray diffraction analysis of the red **fluorescent** protein eqFP611.

L6 ANSWER 4 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
TI Steady-state and time-resolved fluorescence studies indicate an unusual conformation of 2-aminopurine within ATAT and TATA duplex DNA sequences.

L6 ANSWER 5 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
TI A far-red **fluorescent** protein with fast maturation and reduced oligomerization tendency from Entacmaea quadricolor (Anthozoa, Actinaria).

L6 ANSWER 6 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
TI Kinetic analysis of maturation and denaturation of DsRed, a coral-derived red **fluorescent** protein.

L6 ANSWER 7 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN



TI Suitability of enhanced green fluorescent protein as a reporter component for bioassays.

L6 ANSWER 8 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI The structure of the chromophore within DsRed, a red fluorescent protein from coral.

L6 ANSWER 9 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI Novel fluorescent protein from Discosoma coral and its mutants possesses a unique far-red fluorescence.

L6 ANSWER 10 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI **Fluorescent** probing of membrane potential in walled cells: diS-C3(3) assay in *Saccharomyces cerevisiae*.

L6 ANSWER 11 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI Terminal Marking of Triosephosphate Isomerase: Consequences of Deamidation.

L6 ANSWER 12 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI Quenching of intrinsic fluorescence of yeast cytochrome c peroxidase by covalently- and noncovalently-bound quenchers.

L6 ANSWER 13 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI Low temperature absorbance and fluorescence spectroscopy of the photoactive yellow protein from *Ectothiorhodospira halophila*.

L6 ANSWER 14 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI PICOSECOND DECAY KINETICS AND QUANTUM YIELD OF FLUORESCENCE OF THE PHOTOACTIVE YELLOW PROTEIN FROM THE HALOPHILIC PURPLE PHOTOTROPHIC BACTERIUM *ECTOTHIORHODOSPIRA-HALOPHILA*.

L6 ANSWER 15 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI INTERACTION OF MUSCLE AND NON-MUSCLE TROPOMYOSINS WITH DNASE I.

L6 ANSWER 16 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI FLUORESCENCE SPECTROSCOPY OF BENZO-ALPHA-PYRENE DIOL EPOXIDE DNA ADDUCTS CONFORMATION-SPECIFIC **EMISSION** SPECTRA.

L6 ANSWER 17 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI CHARACTERIZATION OF THE FLUORESCENCE OF THE ANTITUMOR AGENT MITOXANTRONE.

L6 ANSWER 18 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI SPECTROFLUOROMETRIC STUDIES OF THE LIPID PROBE NILE RED.

L6 ANSWER 19 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI FLUORESCENCE OF HISTONES H-1 A TYROSINATE-LIKE FLUORESCENCE **EMISSION** IN CERATITIS-CAPITATA H-1 AT NEUTRAL PH VALUES.

L6 ANSWER 20 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
 TI P AMINO BENZAMIDINE AS A **FLUORESCENT** PROBE FOR THE ACTIVE SITE OF SERINE PROTEASES.



=> d 16 ti 21-56

- L6 ANSWER 21 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
TI THE INTERACTION OF HUMAN SERUM ALBUMIN AND HEMOPEXIN WITH PORPHYRINS.
- L6 ANSWER 22 OF 56 MEDLINE on STN  
TI Far-red fluorescent proteins evolved from a blue chromoprotein from Actinia equina.
- L6 ANSWER 23 OF 56 MEDLINE on STN  
TI Photophysics and biological applications of the environment-sensitive fluorophore 6-N,N-dimethylamino-2,3-naphthalimide.
- L6 ANSWER 24 OF 56 MEDLINE on STN  
TI Fluorescence and circular dichroism spectroscopic studies on bovine lactoperoxidase.
- L6 ANSWER 25 OF 56 MEDLINE on STN  
TI Salt-induced folding of alkaline denatured creatine kinase under high pH conditions.
- L6 ANSWER 26 OF 56 MEDLINE on STN  
TI Interaction of tryptophan residues of cytochrome P450<sub>scc</sub> with a highly specific fluorescence quencher, a substrate analogue, compared to acrylamide and iodide.
- L6 ANSWER 27 OF 56 MEDLINE on STN  
TI [Bertalanffy-like fluorescence staining with 3-dimethylamino-6-methoxyacridine].  
Über eine Bertalanffy-analoge Fluorochromierung mit 3-Dimethylamino-6-methoxyacridin.
- L6 ANSWER 28 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Saturated red light-emitting copolymers of poly(aryleneethynylene)s with narrow-band-gap (NBG) units: Synthesis and luminescent properties
- L6 ANSWER 29 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Synthesis and spectral properties of novel water-soluble near-infrared fluorescent indocyanines
- L6 ANSWER 30 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI 7-(2-Methoxycarbonylvinyl)-3-hydroxychromones: new dyes with red shifted dual emission
- L6 ANSWER 31 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI The synthesis and photophysical properties of a novel red-emitting dioxolane-substituted pentacene derivative
- L6 ANSWER 32 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI High-Efficiency Saturated Red Emitting Polymers Derived from Fluorene and Naphthoselenadiazole
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TI Facile synthesis of chlorophyll analog possessing a disulfide bond and formation of self-assembled monolayer on gold surface
- L6 ANSWER 34 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Cd(II) Sensing in Water Using Novel Aromatic Iminodiacetate Based Fluorescent Chemosensors
- L6 ANSWER 35 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Synthesis and Optical and Electroluminescent Properties of Novel

## Conjugated Copolymers Derived from Fluorene and Benzoselenadiazole

- L6 ANSWER 36 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Development of novel cyanine dyes for diode laser induced fluorescence detection
- L6 ANSWER 37 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Synthesis and properties of two polymerizable **fluorescent** monomers
- L6 ANSWER 38 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI RGB **emission** using a dimesitylboryl-bithiophene derivative as a universal host and pentacene derivatives as the red emitters
- L6 ANSWER 39 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Far **red-shifted fluorescent** proteins and their variants from *Heteractis crispis*
- L6 ANSWER 40 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI The first genuine observation of **fluorescent** mononuclear phthalocyanine aggregates
- L6 ANSWER 41 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Probing the donor and acceptor dye assemblies at the galleries of  $\alpha$ -zirconium phosphate
- L6 ANSWER 42 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Natural animal coloration can be determined by a nonfluorescent green **fluorescent** protein homolog
- L6 ANSWER 43 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Synthesis and properties of soluble, **fluorescent** polyesters and polyethers with substituted m-terphenyl segments in the main chain
- L6 ANSWER 44 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI The Formation and Behavior of **Fluorescent** Lewis Acid-Base Exciplexes and Triplexes
- L6 ANSWER 45 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Fluorescence Properties of Benz[f]indole, a Wavelength and Quenching Selective Tryptophan Analog
- L6 ANSWER 46 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Exploring the hydrophobic binding sites of calmodulin with **fluorescent** probes: Dynamic formation of the excimer
- L6 ANSWER 47 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Influence of Chlorination on Chromophores and Fluorophores in Humic Substances
- L6 ANSWER 48 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI An assessment of the usefulness of 5-hydroxytryptophan as an optical probe
- L6 ANSWER 49 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Protonation of free 2,9-bis(p-biphenyl)-1,10-phenanthroline sites in a 56-membered macrocycle and in its ReI and CuI complexes Absorption spectra, luminescence properties, and excited state interactions
- L6 ANSWER 50 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Aggregation state of dehydroergosterol in water and in a model system of membranes
- L6 ANSWER 51 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Fluorescence and Photodimerization Studies of Cyano-Substituted

## Diphenylbutadienes

- L6 ANSWER 52 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Solvent effects on the **fluorescent emission** of some new benzimidazole derivatives
- L6 ANSWER 53 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Toward red-emitting radiation tolerant chromophores
- L6 ANSWER 54 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Chlorotetracycline-induced fluorescence from atherosclerotic plaque with UV laser excitation
- L6 ANSWER 55 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI  $\alpha$ -Tocopherol as a **fluorescent** probe. 1. Its spectroscopic characterization and use as a probe of liposome
- L6 ANSWER 56 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN  
TI Formation of **fluorescent** derivatives of alcohol dehydrogenase and other proteins on UV irradiation

=> d ab bib 1,5,6,8,22,39

L13 HAS NO ANSWERS

'1 5 6 8 22 39 ' IS NOT A VALID SEARCH STATUS KEYWORD

Search status keywords:

NONE ---- Display only the number of postings.

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L13 0 SEA (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM  
AND " RED SHIFTED" AND ANTHOZOAN

=> d 16 ab bib 1,5,6,8,22,39

- L6 ANSWER 1 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
AB Two-photon (TP) excitation (820-1150 nm) and **emission** (280-700 nm) spectra for the **fluorescent** proteins (FPs) ECFP3, EGFP3 and EYFP3 produced in human tumour cells were recorded. TP excitation spectra of pure and highly enriched samples were found to be more differentiated in comparison with their one-photon (OP) spectra. They exhibited more pronounced main and local **maxima**, which coincided among different purity grades within small limits. TP and OP **emission** spectra of pure and enriched samples were identical. However, in crude samples, excitation was slightly blue-shifted and **emission red-shifted**. The data indicate that both OP and TP excitation routes led to the same excited states of these molecules. The **emission** intensity is dependent on the pH of the environment for both types of excitation; the **emission** intensity **maximum** can be recorded in the alkaline range. Reconstitution of **emission** intensity after pH quenching was incomplete, albeit that the respective spectral profiles were identical to those prequenching. When **emission** data were averaged over the whole range of excitation, the resulting **emission** profile and **maximum** coincided with the data generated by optimal excitation. Therefore, out-of-**maximum** excitation, common practice in TP excitation microscopy, can be used for routine application.
- AN 2005:168065 BIOSIS  
DN PREV200500175603  
TI Two-photon excitation and **emission** spectra of the green **fluorescent** protein variants ECFP, EGFP and EYFP.  
AU Spiess, E. [Reprint Author]; Bestvater, F.; Heckel-Pompey, A.; Toth, K.; Hacker, M.; Stobrawa, G.; Feurer, T.; Wotzlaw, C.; Berchner-Pfannschmidt, U.; Porwol, T.; Acker, H.

CS Deutsch Krebsforschungszentrum, D-6900, Heidelberg, Germany  
e.spiess@dkfz-heidelberg.de

SO Journal of Microscopy (Oxford), (March 2005) Vol. 217, No. 3, pp. 200-204.  
print.  
CODEN: JMICAR. ISSN: 0022-2720.

DT Article

LA English

ED Entered STN: 4 May 2005  
Last Updated on STN: 4 May 2005

L6 ANSWER 5 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
AB We performed the biochemical and biophysical characterization of a red  
**fluorescent** protein, eqFP611, from the sea anemone *Entacmaea*  
*quadricolor* cloned in *Escherichia coli*. With an excitation  
**maximum** at 559 nm and an **emission**  
**maximum** at 611 nm, the recombinant protein shows the  
most **red-shifted emission** and the largest  
Stokes shift of all nonmodified proteins in the green **fluorescent**  
protein family. The protein fluoresces with a high quantum yield of 0.45,  
although it resembles the nonfluorescent members of this protein class, as  
inferred from the absence of the key amino acid serine at position 143.  
Fluorescence is constant within the range pH 4-10. Red fluorophore  
maturation reaches a level of 90% after approx 12 h by passing through a  
green intermediate. After complete maturation, only a small fraction of  
the green species (less than 1%) persists. The protein has a reduced  
tendency to oligomerize, as shown by its monomeric appearance in SDS/PAGE  
analysis and single-molecule experiments. However, it forms tetramers at  
higher concentrations in the absence of detergent. Fluorescence  
correlation spectroscopy reveals light-driven transitions between bright  
and dark states on submillisecond and millisecond time scales.  
Applicability of eqFP611 for in vivo labeling in eukaryotic systems was  
shown by expression in a mammalian cell culture.

AN 2002:558092 BIOSIS

DN PREV200200558092

TI A far-red **fluorescent** protein with fast maturation and reduced  
oligomerization tendency from *Entacmaea quadricolor* (Anthozoa, Actinaria).

AU Wiedenmann, Joerg; Schenk, Andreas; Roecker, Carlheinz; Girod, Andreas;  
Spindler, Klaus-Dieter; Nienhaus, G. Ulrich [Reprint author]

CS Department of Physics, University of Illinois at Urbana-Champaign, Urbana,  
IL, 61801, USA  
uli@uiuc.edu

SO Proceedings of the National Academy of Sciences of the United States of  
America, (September 3, 2002) Vol. 99, No. 18, pp. 11646-11651. print.  
CODEN: PNASA6. ISSN: 0027-8424.

DT Article

LA English

OS Genbank-AY130757; EMBL-AY130757; DDBJ-AY130757

ED Entered STN: 30 Oct 2002  
Last Updated on STN: 30 Dec 2002

L6 ANSWER 6 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
AB The red **fluorescent** protein DsRed recently cloned from *Discosoma*  
coral, with its significantly **red-shifted** excitation  
and **emission maxima** (558 and 583 nm,  
respectively), has attracted great interest because of its spectral  
complementation to other **fluorescent** proteins, including the  
green **fluorescent** protein and its enhanced mutant EGFP. We  
demonstrated that the much slower DsRed fluorescence development could be  
described by a three-step kinetic model, in contrast to the fast EGFP  
maturation, which was fitted by a one-step model. At pH below 5.0 DsRed  
fluorescence gradually decreased, and the rate and degree of this  
fluorescence inactivation depended on the pH value. The kinetics of  
fluorescence inactivation under acidic conditions was fitted by a  
two-exponential function where the initial inactivation rate was

proportional to the fourth power of proton concentration. Subsequent DsRed alkalization resulted in partial fluorescence recovery, and the rate and degree of such recovery depended on the incubation time in the acid. Recovery kinetics had a lag-time and was fitted minimally by three exponential functions. The DsRed absorbance and circular dichroism spectra revealed that the fluorescence loss was accompanied by protein denaturation. We developed a kinetic mechanism for DsRed denaturation that includes consecutive conversion of the initial state of the protein, protonated by four hydrogen ions, to the denatured one through three intermediates. The first intermediate still emits fluorescence, and the last one is subjected to irreversible inactivation. Because of tight DsRed tetramerization we have suggested that obligatory protonation of each monomer results in the fluorescence inactivation of the whole tetramer.

AN 2002:149199 BIOSIS  
DN PREV200200149199  
TI Kinetic analysis of maturation and denaturation of DsRed, a coral-derived red fluorescent protein.  
AU Verkhusha, V. V. [Reprint author]; Akovbian, N. A.; Efremenko, E. N.; Varfolomeyev, S. D.; Vrzheschch, P. V.  
CS Center for Molecular Medicine, Lomonosov Moscow State University, Moscow, 119899, Russia  
vrzh@genebee.msu.ru  
SO Biochemistry (Moscow), (December, 2001) Vol. 66, No. 12, pp. 1342-1351. print.  
CODEN: BIORAK. ISSN: 0006-2979.  
DT Article  
LA English  
ED Entered STN: 14 Feb 2002  
Last Updated on STN: 26 Feb 2002

L6 ANSWER 8 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN  
AB DsRed, a brilliantly red fluorescent protein, was recently cloned from Discosoma coral by homology to the green fluorescent protein (GFP) from the jellyfish Aequorea. A core question in the biochemistry of DsRed is the mechanism by which the GFP-like 475-nm excitation and 500-nm emission maxima of immature DsRed are red-shifted to the 558-nm excitation and 583-nm emission maxima of mature DsRed. After digestion of mature DsRed with lysyl endopeptidase, high-resolution mass spectra of the purified chromophore-bearing peptide reveal that some of the molecules have lost 2 Da relative to the peptide analogously prepared from a mutant, K83R, that stays green. Tandem mass spectrometry indicates that the bond between the alpha-carbon and nitrogen of Gln-66 has been dehydrogenated in DsRed, extending the GFP chromophore by forming -CdbdN-CdbdO at the 2-position of the imidazolidinone. This acylimine substituent quantitatively accounts for the red shift according to quantum mechanical calculations. Reversible hydration of the CdbdN bond in the acylimine would explain why denaturation shifts mature DsRed back to a GFP-like absorbance. The CdbdN bond hydrolyses upon boiling, explaining why DsRed shows two fragment bands on SDS/PAGE. This assay suggests that conversion from green to red chromophores remains incomplete even after prolonged aging.

AN 2001:12180 BIOSIS  
DN PREV200100012180  
TI The structure of the chromophore within DsRed, a red fluorescent protein from coral.  
AU Gross, Larry A.; Baird, Geoffrey S.; Hoffman, Ross C.; Baldrige, Kim K.; Tsien, Roger Y. [Reprint author]  
CS University of California, San Diego, 9500 Gilman Drive, 310 Cellular and Molecular Medicine West 0647, La Jolla, CA, 92093-0647, USA  
SO Proceedings of the National Academy of Sciences of the United States of America, (October 24, 2000) Vol. 97, No. 22, pp. 11990-11995. print.  
CODEN: PNASA6. ISSN: 0027-8424.



DT Article  
LA English  
ED Entered STN: 27 Dec 2000  
Last Updated on STN: 27 Dec 2000

L6 ANSWER 22 OF 56 MEDLINE on STN

AB Proteins of the GFP (green fluorescent protein) family demonstrate a great spectral and phylogenetic diversity. However, there is still an intense demand for **red-shifted** GFP-like proteins in both basic and applied science. To obtain GFP-like **chromoproteins** with **red-shifted** absorption, we performed a broad search in blue-coloured Anthozoa species. We revealed specimens of *Actinia equina* (beadlet anemone) exhibiting a bright blue circle band at the edge of the basal disc. A novel blue **chromoprotein**, aeCP597, with an absorption **maximum** at 597 nm determining the coloration of the anemone basal disk was cloned. AeCP597 carries a chromophore chemically identical with that of the well-studied DsRed (red fluorescent protein from *Discosoma* sp.). Thus a strong 42-nm bathochromic shift of aeCP597 absorption compared with DsRed is determined by peculiarities of chromophore environment. Site-directed and random mutagenesis of aeCP597 resulted in far-red **fluorescent** mutants with **emission maxima** at up to 663 nm. The most bright and stable mutant AQ143 possessed excitation and **emission maxima** at 595 and 655 nm respectively. Thus aeCP597 and its **fluorescent** mutants set a new record of **red-shifted** absorption and **emission maxima** among GFP-like proteins.

AN 2005640113 IN-PROCESS

DN PubMed ID: 16164420

TI Far-red **fluorescent** proteins evolved from a blue **chromoprotein** from *Actinia equina*.

AU Shkrob Maria A; Yanushevich Yurii G; Chudakov Dmitriy M; Gurskaya Nadya G; Labas Yulii A; Poponov Sergey Y; Mudrik Nikolay N; Lukyanov Sergey; Lukyanov Konstantin A

CS Shemiakin-Ovchinnikov Institute of Bioorganic Chemistry, Miklukho-Maklaya 16/10, 117997 Moscow, Russia.

SO Biochemical journal, (2005 Dec 15) 392 (Pt 3) 649-54.  
Journal code: 2984726R. ISSN: 1470-8728.

CY England: United Kingdom

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS NONMEDLINE; IN-DATA-REVIEW; IN-PROCESS; NONINDEXED; Priority Journals

ED Entered STN: 20051203

Last Updated on STN: 20051203

L6 ANSWER 39 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN

AB Nucleic acid compns. encoding Stichodactylidae **chromoproteins** and **fluorescent** mutants thereof, as well as the polypeptide compns. encoded by the same, are provided. The proteins of interest are proteins that are colored and/or **fluorescent**, where this feature arises from the interaction of two or more residues of the protein. Also of interest are proteins that are substantially similar to, or mutants of, the above specific proteins, including non-aggregating mutants and mutants with modulated oligomerization characteristics as compared to wild type. Thus, the two wild-type **chromoprotein** isoforms from *Heteractis crispa* exhibit a strong **emission max.** at .apprx.580-640 nm. Site-specific mutagenesis of the Cys-148 residue to serine dramatically increases the quantum yield of red fluorescence as compared to the wild-type protein, and further random mutagenesis (A2S, T36A, C143S, L173H, P201L, K204E) yielded an even brighter mutant. A single mutation, L126H, may be responsible for modifying the oligomeric state of the protein from tetrameric to dimeric. Also provided are fragments of the nucleic acids and the peptides encoded

thereby, as well as antibodies to the subject proteins and transgenic cells and organisms. The subject protein and nucleic acid compns. find use in a variety of different applications. Finally, kits for use in such applications, e.g., that include the subject nucleic acid compns., are provided.

AN 2002:293685 CAPLUS

DN 136:321045

TI Far **red-shifted fluorescent** proteins and their variants from *Heteractis crispis*

IN Lukyanov, Sergey Anatolievich; Fradkov, Arcady Fedorovich; Lukyanov, Konstantin Anatolievich; Gurskaya, Nadezda Georgievna

PA Clontech Laboratories, Inc., USA

SO PCT Int. Appl., 81 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 17

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002030965	A2	20020418	WO 2001-US32080	20011012
	WO 2002030965	A3	20030605		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2425318	AA	20020418	CA 2001-2425318	20011012
	AU 2002011722	A5	20020422	AU 2002-11722	20011012
	EP 1334122	A2	20030813	EP 2001-979797	20011012
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2004532606	T2	20041028	JP 2002-534350	20011012
PRAI	US 2000-240018P	P	20001012		
	US 2001-306131P	P	20010716		
	WO 2001-US32080	W	20011012		